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NEWS	4	DEC 08	INPADOC: Legal Status data reloaded
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NEWS	7	OCT 21	BIOSIS file reloaded and enhanced
NEWS	8	OCT 28	BIOSIS file segment of TOXCENTER reloaded and enhanced
NEWS	9	NOV 24	MSDS-CCOHS file reloaded
NEWS	10	DEC 08	CABA reloaded with left truncation
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NEWS	16	DEC 19	CROPU no longer updated; subscriber discount no longer available
NEWS	17	DEC 22	Additional INPI reactions and pre-1907 documents added to CAS databases
NEWS	18	DEC 22	IFIPAT/IFIUDB/IFICDB reloaded with new data and search fields
NEWS	19	DEC 22	ABI-INFORM now available on STN
NEWS	20	JAN 27	Source of Registration (SR) information in REGISTRY updated and searchable
NEWS	21	JAN 27	A new search aid, the Company Name Thesaurus, available in CA/CAPLUS
NEWS	22	FEB 05	German (DE) application and patent publication number format changes
NEWS EXPRESS			DECEMBER 28 CURRENT WINDOWS VERSION IS V7.00, CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP), AND CURRENT DISCOVER FILE IS DATED 23 SEPTEMBER 2003
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FILE 'HOME' ENTERED AT 13:35:55 ON 13 FEB 2004

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COST IN U.S. DOLLARS	SINCE FILE	TOTAL
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FULL ESTIMATED COST	0.21	0.21

FILE 'MEDLINE' ENTERED AT 13:36:29 ON 13 FEB 2004

FILE 'USPATFULL' ENTERED AT 13:36:29 ON 13 FEB 2004  
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=> s alzheimer's disease  
MISMATCHED QUOTE 'ALZHEIMER'S'  
Quotation marks (or apostrophes) must be used in pairs,  
one before and one after the expression you are setting  
off or masking.

=> s alzheimer disease  
L1 92358 ALZHEIMER DISEASE

=> s l1 and insulin  
L2 1284 L1 AND INSULIN

=> s insulin ailments  
L3 0 INSULIN AILMENTS

=> s insulin () disease  
L4 18 INSULIN (W) DISEASE

=> s l4 and l1  
L5 1 L4 AND L1

=> d l5 ti abs ibib tot

L5 ANSWER 1 OF 1 USPATFULL on STN  
TI 207 human secreted proteins  
AB The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:258639 USPATFULL  
TITLE: 207 human secreted proteins  
INVENTOR(S): Ni, Jian, Germantown, MD, UNITED STATES  
Ebner, Reinhard, Gaithersburg, MD, UNITED STATES

LaFleur, David W., Washington, DC, UNITED STATES  
 Moore, Paul A., Germantown, MD, UNITED STATES  
 Olsen, Henrik S., Gaithersburg, MD, UNITED STATES  
 Rosen, Craig A., Laytonsville, MD, UNITED STATES  
 Ruben, Steven M., Olney, MD, UNITED STATES  
 Soppet, Daniel R., Centreville, VA, UNITED STATES  
 Young, Paul E., Gaithersburg, MD, UNITED STATES  
 Shi, Yanggu, Gaithersburg, MD, UNITED STATES  
 Florence, Kimberly A., Rockville, MD, UNITED STATES  
 Wei, Ying-Fei, Berkeley, CA, UNITED STATES  
 Florence, Charles, Rockville, MD, UNITED STATES  
 Hu, Jing-Shan, Mountain View, CA, UNITED STATES  
 Li, Yi, Sunnyvale, CA, UNITED STATES  
 Kyaw, Hla, Frederick, MD, UNITED STATES  
 Fischer, Carrie L., Burke, VA, UNITED STATES  
 Ferrie, Ann M., Painted Post, NY, UNITED STATES  
 Fan, Ping, Potomac, MD, UNITED STATES  
 Feng, Ping, Gaithersburg, MD, UNITED STATES  
 Endress, Gregory A., Florence, MA, UNITED STATES  
 Dillon, Patrick J., Carlsbad, CA, UNITED STATES  
 Carter, Kenneth C., North Potomac, MD, UNITED STATES  
 Brewer, Laurie A., St. Paul, MN, UNITED STATES  
 Yu, Guo-Liang, Berkeley, CA, UNITED STATES  
 Zeng, Zhizhen, Lansdale, PA, UNITED STATES  
 Greene, John M., Gaithersburg, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003181692	A1	20030925
APPLICATION INFO.:	US 2001-933767	A1	20010822 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2001-US5614, filed on 21 Feb 2001, PENDING Continuation-in-part of Ser. No. US 1998-205258, filed on 4 Dec 1998, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-184836P	20000224 (60)
	US 2000-193170P	20000329 (60)
	US 1997-48885P	19970606 (60)
	US 1997-49375P	19970606 (60)
	US 1997-48881P	19970606 (60)
	US 1997-48880P	19970606 (60)
	US 1997-48896P	19970606 (60)
	US 1997-49020P	19970606 (60)
	US 1997-48876P	19970606 (60)
	US 1997-48895P	19970606 (60)
	US 1997-48884P	19970606 (60)
	US 1997-48894P	19970606 (60)
	US 1997-48971P	19970606 (60)
	US 1997-48964P	19970606 (60)
	US 1997-48882P	19970606 (60)
	US 1997-48899P	19970606 (60)
	US 1997-48893P	19970606 (60)
	US 1997-48900P	19970606 (60)
	US 1997-48901P	19970606 (60)
	US 1997-48892P	19970606 (60)
	US 1997-48915P	19970606 (60)
	US 1997-49019P	19970606 (60)
	US 1997-48970P	19970606 (60)
	US 1997-48972P	19970606 (60)
	US 1997-48916P	19970606 (60)
	US 1997-49373P	19970606 (60)
	US 1997-48875P	19970606 (60)
	US 1997-49374P	19970606 (60)

US 1997-48917P	19970606 (60)
US 1997-48949P	19970606 (60)
US 1997-48974P	19970606 (60)
US 1997-48883P	19970606 (60)
US 1997-48897P	19970606 (60)
US 1997-48898P	19970606 (60)
US 1997-48962P	19970606 (60)
US 1997-48963P	19970606 (60)
US 1997-48877P	19970606 (60)
US 1997-48878P	19970606 (60)
US 1997-57645P	19970905 (60)
US 1997-57642P	19970905 (60)
US 1997-57668P	19970905 (60)
US 1997-57635P	19970905 (60)
US 1997-57627P	19970905 (60)
US 1997-57667P	19970905 (60)
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US 1997-57776P	19970905 (60)
US 1997-57778P	19970905 (60)
US 1997-57629P	19970905 (60)
US 1997-57628P	19970905 (60)
US 1997-57777P	19970905 (60)
US 1997-57634P	19970905 (60)
US 1997-70923P	19971218 (60)
US 1998-92921P	19980715 (60)
US 1998-94657P	19980730 (60)
US 1997-70923P	19971218 (60)
US 1998-92921P	19980715 (60)
US 1998-94657P	19980730 (60)

DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION  
LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
ROCKVILLE, MD, 20850  
NUMBER OF CLAIMS: 23  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 10 Drawing Page(s)  
LINE COUNT: 32746  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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(FILE 'HOME' ENTERED AT 13:35:55 ON 13 FEB 2004)

FILE 'MEDLINE, USPATFULL, DGENE, EMBASE, WPIDS, BIOSIS' ENTERED AT  
13:36:29 ON 13 FEB 2004

L1 92358 S ALZHEIMER DISEASE  
L2 1284 S L1 AND INSULIN  
L3 0 S INSULIN AILMENTS  
L4 18 S INSULIN ( ) DISEASE  
L5 1 S L4 AND L1

=> s glucose therapy and l1  
L6 0 GLUCOSE THERAPY AND L1

=> s glucose therapy  
L7 159 GLUCOSE THERAPY

=> s insulin therapy\  
L8 11261 INSULIN THERAPY\  
L9 10 L8 AND L1

=> s l8 and l1  
L9 10 L8 AND L1

=> s l7 and l1  
L10 0 L7 AND L1

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L9 ANSWER 1 OF 10 USPATFULL on STN

TI Synthetic immunogenic but non-deposit-forming polypeptides and peptides homologous to amyloid beta, prion protein, amylin, alpha-synuclein, or polyglutamine repeats for induction of an immune response thereto  
AB The present invention relates to immunogenic but non-depositing-forming polypeptides or peptides homologous to amyloid  $\beta$ , prion, amylin or  $\alpha$ -synuclein which can be used alone or conjugated to an immunostimulatory molecule in an immunizing composition for inducing an immune response to amyloid  $\beta$  peptides and amyloid deposits, to prion protein and prion deposits, to amylin and amylin deposits, to  $\alpha$ -synuclein and deposits containing  $\alpha$ -synuclein, or to polyglutamine repeats and deposits of proteins containing polyglutamine repeats. Described are also antibodies directed against such peptides, their generation, and their use in methods of passive immunization to such peptides and deposits.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:238400 USPATFULL

TITLE: Synthetic immunogenic but non-deposit-forming polypeptides and peptides homologous to amyloid beta, prion protein, amylin, alpha-synuclein, or polyglutamine repeats for induction of an immune response thereto

INVENTOR(S): Frangione, Blas, New York, NY, UNITED STATES  
Wisniewski, Thomas, Statent Island, NY, UNITED STATES  
Sigurdsson, Einar M., New York, NY, UNITED STATES

PATENT ASSIGNEE(S): NEW YORK UNIVERSITY (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003166558	A1	20030904
APPLICATION INFO.:	US 2002-301488	A1	20021121 (10)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2001-331801P	20011121 (60)

DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION  
LEGAL REPRESENTATIVE: DARBY & DARBY P.C., Post Office Box 5257, New York, NY,  
10150-5257  
NUMBER OF CLAIMS: 115  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 33 Drawing Page(s)  
LINE COUNT: 4966  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 2 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:30987 USPATFULL  
TITLE: Therapeutic compositions  
INVENTOR(S): Veech, Richard L., Rockville, MD, UNITED STATES  
PATENT ASSIGNEE(S): BTG International Limited (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003022937	A1	20030130
APPLICATION INFO.:	US 2002-153873	A1	20020524 (10)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 2001-843694, filed on 30 Apr 2001, ABANDONED Continuation of Ser. No. US 1999-397100, filed on 16 Sep 1999, GRANTED, Pat. No. US 6323237 Continuation of Ser. No. WO 1998-US5072, filed on 17 Mar 1998, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	NIXON & VANDERHUYE P.C., 8th Floor, 1100 North Glebe Road, Arlington, VA, 22201	
NUMBER OF CLAIMS:	31	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Page(s)	
LINE COUNT:	1883	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 3 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant

states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:215082 USPATFULL  
TITLE: Therapeutic compositions  
INVENTOR(S): Veech, Richard L., Rockville, MD, United States  
PATENT ASSIGNEE(S): BTG International Limited, London, United Kingdom  
(non-U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6323237	B1	20011127
APPLICATION INFO.:	US 1999-397100		19990916 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 1998-US5072, filed on 17 Mar 1998		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Reamer, James H.	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye	
NUMBER OF CLAIMS:	19	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 2 Drawing Page(s)	
LINE COUNT:	2039	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 4 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:205943 USPATFULL  
TITLE: Therapeutic compositions  
INVENTOR(S): Veech, Richard L., Rockville, MD, United States  
PATENT ASSIGNEE(S): BTG International Limited (U.S. corporation)

NUMBER	KIND	DATE
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PATENT INFORMATION: US 2001041736 A1 20011115  
 APPLICATION INFO.: US 2001-843694 A1 20010430 (9)  
 RELATED APPLN. INFO.: Continuation of Ser. No. US 1999-397100, filed on 16  
 Sep 1999, PENDING Continuation of Ser. No. WO  
 1998-US5072, filed on 17 Mar 1998, UNKNOWN

	NUMBER	DATE
	-----	-----
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye P.C., 8th Floor, 1100 N. Glebe Rd., Arlington, VA, 22201	
NUMBER OF CLAIMS:	31	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Page(s)	
LINE COUNT:	1889	
CAS INDEXING IS AVAILABLE FOR THIS PATENT.		

L9 ANSWER 5 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:202234 USPATFULL  
 TITLE: Therapeutic compositions  
 INVENTOR(S): Veech, Richard Lewis, Rockville, MD, United States  
 PATENT ASSIGNEE(S): BTG International Limited, London, United Kingdom  
 (non-U.S. corporation)

	NUMBER	KIND	DATE
	-----	-----	-----
PATENT INFORMATION:	US 6316038	B1	20011113
APPLICATION INFO.:	US 1999-397109		19990916 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. WO 1998-GB5072, filed on 17 Mar 1998		

	NUMBER	DATE
	-----	-----
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Reamer, James H.	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye	
NUMBER OF CLAIMS:	2	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	1 Drawing Figure(s); 1 Drawing Page(s)	
LINE COUNT:	1821	
CAS INDEXING IS AVAILABLE FOR THIS PATENT.		



L9 ANSWER 6 OF 10 USPATFULL on STN

TI Therapeutic compositions (II)

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:134247 USPATFULL

TITLE: Therapeutic compositions (II)

INVENTOR(S): Veech, Richard Lewis, Rockville, MD, United States

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2001014696	A1	20010816
APPLICATION INFO.:	US 2001-799124	A1	20010306 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. WO 1999-US21015, filed on 15 Sep 1999, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1998-100371P	19980915 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye, Eighth Floor, 1100 North Glebe Road, Arlington, VA, 22201-4714	
NUMBER OF CLAIMS:	18	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	1 Drawing Page(s)	
LINE COUNT:	1376	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 7 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:44413 USPATFULL

TITLE: Therapeutic compositions

INVENTOR(S): Veech, Richard L., Rockville, MD, United States  
PATENT ASSIGNEE(S): BTG International Limited, London, United Kingdom  
(non-U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6207856	B1	20010327
APPLICATION INFO.:	US 2000-630007		20000731 (9)
RELATED APPLN. INFO.:	Division of Ser. No. US 1999-397100, filed on 16 Sep 1999 Continuation of Ser. No. WO 1997-US9805072, filed on 17 Mar 1997		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Reamer, James H.	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye	
NUMBER OF CLAIMS:	4	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 2 Drawing Page(s)	
LINE COUNT:	1949	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 8 OF 10 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.  
on STN

TI Butyrylcholinesterase K variant on chromosome 3 q is associated with Type II diabetes in white Caucasian subjects.

AB Aims/hypothesis. To determine the association of three genes associated with Alzheimer's disease - butyrylcholinesterase (BcHE) on chromosome 3 q,  $\alpha 2$  macroglobulin ( $\alpha 2$ M) on chromosome 12p and apolipoprotein E (ApoE) on chromosome 19 q -with Type II (non-insulin-dependent) diabetes mellitus. Methods. Frequencies of BcHE K variant,  $\alpha 2$ M insertion and/or deletion polymorphism, the ApoE common polymorphisms and promoter variants at ApoE-491 and -291, were examined by fluorescent RFLP in DNA from 276 United Kingdom Prospective Diabetes Study Type II diabetic subjects and 351 non-diabetic subjects from the Diabetes In Families study. Genetic data in diabetic subjects was analysed in relation to clinical characteristics and islet function as assessed by the requirement for **insulin therapy** 6 years after randomisation.

Results. The BcHE K variant allele was more common among Type II diabetic subjects (D) than non-diabetic subjects (ND) (22.8% D vs 15.8% ND;  $p = 0.00017$ ). Subjects homozygous for the variant were more frequent in the diabetic group (5.8% D vs 2.6% ND;  $p = 0.00056$ ). The K variant allele frequency was not associated with a requirement for **insulin therapy** (29.0% insulin-requiring vs 21.8% non-insulin-requiring;  $p = 0.121$ ). There were no associations of  $\alpha 2$ M and ApoE polymorphisms or ApoE promoter variants with clinical characteristics or insulin requirement in diabetic subjects. There were differences in total cholesterol ( $p = 0.0005$ ) and LDL-cholesterol ( $p = 0.0009$ ) among non-diabetic subjects in relation to ApoE-491 genotypes.

Conclusion/interpretation. The association of the BcHE gene (3q26) with Type II diabetes could be related to an identified susceptibility locus on chromosome 3q27 but appears to be independent of islet function. The absence of diabetes-specific associations with  $\alpha 2$ M, ApoE or ApoE promoter variants suggest that these are not important in the onset of hyperglycaemia.

ACCESSION NUMBER: 2002005139 EMBASE

TITLE: Butyrylcholinesterase K variant on chromosome 3 q is associated with Type II diabetes in white Caucasian subjects.

AUTHOR: Hashim Y.; Shepherd D.; Wiltshire S.; Holman R.R.; Levy J.C.; Clark A.; Cull C.A.

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID:sssptal653hxp

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2		"Ask CAS" for self-help around the clock
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NEWS	5	SEP 29	DISSABS now available on STN
NEWS	6	OCT 10	PCTFULL: Two new display fields added
NEWS	7	OCT 21	BIOSIS file reloaded and enhanced
NEWS	8	OCT 28	BIOSIS file segment of TOXCENTER reloaded and enhanced
NEWS	9	NOV 24	MSDS-CCOHS file reloaded
NEWS	10	DEC 08	CABA reloaded with left truncation
NEWS	11	DEC 08	IMS file names changed
NEWS	12	DEC 09	Experimental property data collected by CAS now available in REGISTRY
NEWS	13	DEC 09	STN Entry Date available for display in REGISTRY and CA/CAPLUS
NEWS	14	DEC 17	DGENE: Two new display fields added
NEWS	15	DEC 18	BIOTECHNO no longer updated
NEWS	16	DEC 19	CROPU no longer updated; subscriber discount no longer available
NEWS	17	DEC 22	Additional INPI reactions and pre-1907 documents added to CAS databases
NEWS	18	DEC 22	IFIPAT/IFIUDB/IFICDB reloaded with new data and search fields
NEWS	19	DEC 22	ABI-INFORM now available on STN
NEWS	20	JAN 27	Source of Registration (SR) information in REGISTRY updated and searchable
NEWS	21	JAN 27	A new search aid, the Company Name Thesaurus, available in CA/CAPLUS
NEWS	22	FEB 05	German (DE) application and patent publication number format changes
NEWS EXPRESS			DECEMBER 28 CURRENT WINDOWS VERSION IS V7.00, CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP), AND CURRENT DISCOVER FILE IS DATED 23 SEPTEMBER 2003
NEWS HOURS			STN Operating Hours Plus Help Desk Availability
NEWS INTER			General Internet Information
NEWS LOGIN			Welcome Banner and News Items
NEWS PHONE			Direct Dial and Telecommunication Network Access to STN
NEWS WWW			CAS World Wide Web Site (general information)

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FILE 'HOME' ENTERED AT 13:35:55 ON 13 FEB 2004

=> file medline, uspatful, dgene, embase, wpids, biosis		
COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'MEDLINE' ENTERED AT 13:36:29 ON 13 FEB 2004

FILE 'USPATFULL' ENTERED AT 13:36:29 ON 13 FEB 2004  
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FILE 'BIOSIS' ENTERED AT 13:36:29 ON 13 FEB 2004  
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=> s alzheimer's disease  
MISMATCHED QUOTE 'ALZHEIMER'S'  
Quotation marks (or apostrophes) must be used in pairs,  
one before and one after the expression you are setting  
off or masking.

=> s alzheimer disease  
L1 92358 ALZHEIMER DISEASE

=> s l1 and insulin  
L2 1284 L1 AND INSULIN

=> s insulin ailments  
L3 0 INSULIN AILMENTS

=> s insulin () disease  
L4 18 INSULIN (W) DISEASE

=> s l4 and l1  
L5 1 L4 AND L1

=> d l5 ti abs ibib tot

L5 ANSWER 1 OF 1 USPATFULL on STN  
TI 207 human secreted proteins  
AB The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:258639 USPATFULL  
TITLE: 207 human secreted proteins  
INVENTOR(S): Ni, Jian, Germantown, MD, UNITED STATES  
Ebner, Reinhard, Gaithersburg, MD, UNITED STATES

LaFleur, David W., Washington, DC, UNITED STATES  
 Moore, Paul A., Germantown, MD, UNITED STATES  
 Olsen, Henrik S., Gaithersburg, MD, UNITED STATES  
 Rosen, Craig A., Laytonsville, MD, UNITED STATES  
 Ruben, Steven M., Olney, MD, UNITED STATES  
 Soppet, Daniel R., Centreville, VA, UNITED STATES  
 Young, Paul E., Gaithersburg, MD, UNITED STATES  
 Shi, Yanggu, Gaithersburg, MD, UNITED STATES  
 Florence, Kimberly A., Rockville, MD, UNITED STATES  
 Wei, Ying-Fei, Berkeley, CA, UNITED STATES  
 Florence, Charles, Rockville, MD, UNITED STATES  
 Hu, Jing-Shan, Mountain View, CA, UNITED STATES  
 Li, Yi, Sunnyvale, CA, UNITED STATES  
 Kyaw, Hla, Frederick, MD, UNITED STATES  
 Fischer, Carrie L., Burke, VA, UNITED STATES  
 Ferrie, Ann M., Painted Post, NY, UNITED STATES  
 Fan, Ping, Potomac, MD, UNITED STATES  
 Feng, Ping, Gaithersburg, MD, UNITED STATES  
 Endress, Gregory A., Florence, MA, UNITED STATES  
 Dillon, Patrick J., Carlsbad, CA, UNITED STATES  
 Carter, Kenneth C., North Potomac, MD, UNITED STATES  
 Brewer, Laurie A., St. Paul, MN, UNITED STATES  
 Yu, Guo-Liang, Berkeley, CA, UNITED STATES  
 Zeng, Zhizhen, Lansdale, PA, UNITED STATES  
 Greene, John M., Gaithersburg, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003181692	A1	20030925
APPLICATION INFO.:	US 2001-933767	A1	20010822 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2001-US5614, filed on 21 Feb 2001, PENDING Continuation-in-part of Ser. No. US 1998-205258, filed on 4 Dec 1998, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-184836P	20000224 (60)
	US 2000-193170P	20000329 (60)
	US 1997-48885P	19970606 (60)
	US 1997-49375P	19970606 (60)
	US 1997-48881P	19970606 (60)
	US 1997-48880P	19970606 (60)
	US 1997-48896P	19970606 (60)
	US 1997-49020P	19970606 (60)
	US 1997-48876P	19970606 (60)
	US 1997-48895P	19970606 (60)
	US 1997-48884P	19970606 (60)
	US 1997-48894P	19970606 (60)
	US 1997-48971P	19970606 (60)
	US 1997-48964P	19970606 (60)
	US 1997-48882P	19970606 (60)
	US 1997-48899P	19970606 (60)
	US 1997-48893P	19970606 (60)
	US 1997-48900P	19970606 (60)
	US 1997-48901P	19970606 (60)
	US 1997-48892P	19970606 (60)
	US 1997-48915P	19970606 (60)
	US 1997-49019P	19970606 (60)
	US 1997-48970P	19970606 (60)
	US 1997-48972P	19970606 (60)
	US 1997-48916P	19970606 (60)
	US 1997-49373P	19970606 (60)
	US 1997-48875P	19970606 (60)
	US 1997-49374P	19970606 (60)

US 1997-48917P	19970606 (60)
US 1997-48949P	19970606 (60)
US 1997-48974P	19970606 (60)
US 1997-48883P	19970606 (60)
US 1997-48897P	19970606 (60)
US 1997-48898P	19970606 (60)
US 1997-48962P	19970606 (60)
US 1997-48963P	19970606 (60)
US 1997-48877P	19970606 (60)
US 1997-48878P	19970606 (60)
US 1997-57645P	19970905 (60)
US 1997-57642P	19970905 (60)
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US 1997-57635P	19970905 (60)
US 1997-57627P	19970905 (60)
US 1997-57667P	19970905 (60)
US 1997-57666P	19970905 (60)
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US 1997-57643P	19970905 (60)
US 1997-57769P	19970905 (60)
US 1997-57763P	19970905 (60)
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US 1997-57584P	19970905 (60)
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US 1997-57778P	19970905 (60)
US 1997-57629P	19970905 (60)
US 1997-57628P	19970905 (60)
US 1997-57777P	19970905 (60)
US 1997-57634P	19970905 (60)
US 1997-70923P	19971218 (60)
US 1998-92921P	19980715 (60)
US 1998-94657P	19980730 (60)
US 1997-70923P	19971218 (60)
US 1998-92921P	19980715 (60)
US 1998-94657P	19980730 (60)

DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION  
LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
ROCKVILLE, MD, 20850  
NUMBER OF CLAIMS: 23  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 10 Drawing Page(s)  
LINE COUNT: 32746  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

=> d his

(FILE 'HOME' ENTERED AT 13:35:55 ON 13 FEB 2004)

FILE 'MEDLINE, USPATFULL, DGENE, EMBASE, WPIDS, BIOSIS' ENTERED AT  
13:36:29 ON 13 FEB 2004

L1 92358 S ALZHEIMER DISEASE  
L2 1284 S L1 AND INSULIN  
L3 0 S INSULIN AILMENTS  
L4 18 S INSULIN ( ) DISEASE  
L5 1 S L4 AND L1

=> s glucose therapy and l1  
L6 0 GLUCOSE THERAPY AND L1

=> s glucose therapy  
L7 159 GLUCOSE THERAPY

=> s insulin therapy\  
L8 11261 INSULIN THERAPY\  
L9

=> s l8 and l1  
L9 10 L8 AND L1

=> s l7 and l1  
L10 0 L7 AND L1

=> d l9 ti abs ibib tot

L9 ANSWER 1 OF 10 USPATFULL on STN

TI Synthetic immunogenic but non-deposit-forming polypeptides and peptides homologous to amyloid beta, prion protein, amylin, alpha-synuclein, or polyglutamine repeats for induction of an immune response thereto  
AB The present invention relates to immunogenic but non-depositing-forming polypeptides or peptides homologous to amyloid  $\beta$ , prion, amylin or  $\alpha$ -synuclein which can be used alone or conjugated to an immunostimulatory molecule in an immunizing composition for inducing an immune response to amyloid  $\beta$  peptides and amyloid deposits, to prion protein and prion deposits, to amylin and amylin deposits, to  $\alpha$ -synuclein and deposits containing  $\alpha$ -synuclein, or to polyglutamine repeats and deposits of proteins containing polyglutamine repeats. Described are also antibodies directed against such peptides, their generation, and their use in methods of passive immunization to such peptides and deposits.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:238400 USPATFULL

TITLE: Synthetic immunogenic but non-deposit-forming polypeptides and peptides homologous to amyloid beta, prion protein, amylin, alpha-synuclein, or polyglutamine repeats for induction of an immune response thereto

INVENTOR(S): Frangione, Blas, New York, NY, UNITED STATES  
Wisniewski, Thomas, Statent Island, NY, UNITED STATES  
Sigurdsson, Einar M., New York, NY, UNITED STATES

PATENT ASSIGNEE(S): NEW YORK UNIVERSITY (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003166558	A1	20030904
APPLICATION INFO.:	US 2002-301488	A1	20021121 (10)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2001-331801P	20011121 (60)

DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION  
LEGAL REPRESENTATIVE: DARBY & DARBY P.C., Post Office Box 5257, New York, NY,  
10150-5257  
NUMBER OF CLAIMS: 115  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 33 Drawing Page(s)  
LINE COUNT: 4966  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 2 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:30987 USPATFULL  
TITLE: Therapeutic compositions  
INVENTOR(S): Veech, Richard L., Rockville, MD, UNITED STATES  
PATENT ASSIGNEE(S): BTG International Limited (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003022937	A1	20030130
APPLICATION INFO.:	US 2002-153873	A1	20020524 (10)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 2001-843694, filed on 30 Apr 2001, ABANDONED Continuation of Ser. No. US 1999-397100, filed on 16 Sep 1999, GRANTED, Pat. No. US 6323237 Continuation of Ser. No. WO 1998-US5072, filed on 17 Mar 1998, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	NIXON & VANDERHYE P.C., 8th Floor, 1100 North Glebe Road, Arlington, VA, 22201	
NUMBER OF CLAIMS:	31	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Page(s)	
LINE COUNT:	1883	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 3 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant



states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:215082 USPATFULL  
TITLE: Therapeutic compositions  
INVENTOR(S): Veech, Richard L., Rockville, MD, United States  
PATENT ASSIGNEE(S): BTG International Limited, London, United Kingdom  
(non-U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6323237	B1	20011127
APPLICATION INFO.:	US 1999-397100		19990916 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 1998-US5072, filed on 17 Mar 1998		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Reamer, James H.	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye	
NUMBER OF CLAIMS:	19	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 2 Drawing Page(s)	
LINE COUNT:	2039	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 4 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:205943 USPATFULL  
TITLE: Therapeutic compositions  
INVENTOR(S): Veech, Richard L., Rockville, MD, United States  
PATENT ASSIGNEE(S): BTG International Limited (U.S. corporation)

NUMBER	KIND	DATE
-----	-----	-----

PATENT INFORMATION: US 2001041736 A1 20011115  
APPLICATION INFO.: US 2001-843694 A1 20010430 (9)  
RELATED APPLN. INFO.: Continuation of Ser. No. US 1999-397100, filed on 16  
Sep 1999, PENDING Continuation of Ser. No. WO  
1998-US5072, filed on 17 Mar 1998, UNKNOWN

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye P.C., 8th Floor, 1100 N. Glebe Rd., Arlington, VA, 22201	
NUMBER OF CLAIMS:	31	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Page(s)	
LINE COUNT:	1889	
CAS INDEXING IS AVAILABLE FOR THIS PATENT.		

L9 ANSWER 5 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:202234 USPATFULL  
TITLE: Therapeutic compositions  
INVENTOR(S): Veech, Richard Lewis, Rockville, MD, United States  
PATENT ASSIGNEE(S): BTG International Limited, London, United Kingdom  
(non-U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6316038	B1	20011113
APPLICATION INFO.:	US 1999-397109		19990916 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. WO 1998-GB5072, filed on 17 Mar 1998		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Reamer, James H.	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye	
NUMBER OF CLAIMS:	2	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	1 Drawing Figure(s); 1 Drawing Page(s)	
LINE COUNT:	1821	
CAS INDEXING IS AVAILABLE FOR THIS PATENT.		

L9 ANSWER 6 OF 10 USPATFULL on STN

TI Therapeutic compositions (II)

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:134247 USPATFULL

TITLE: Therapeutic compositions (II)

INVENTOR(S): Veech, Richard Lewis, Rockville, MD, United States

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2001014696	A1	20010816
APPLICATION INFO.:	US 2001-799124	A1	20010306 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. WO 1999-US21015, filed on 15 Sep 1999, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1998-100371P	19980915 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye, Eighth Floor, 1100 North Glebe Road, Arlington, VA, 22201-4714	
NUMBER OF CLAIMS:	18	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	1 Drawing Page(s)	
LINE COUNT:	1376	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 7 OF 10 USPATFULL on STN

TI Therapeutic compositions

AB Compositions comprising ketone bodies and/or their metabolic precursors are provided that are suitable for administration to humans and animals and which have the properties of, inter alia, (i) increasing cardiac efficiency, particularly efficiency in use of glucose, (ii) for providing energy source, particularly in diabetes and insulin resistant states and (iii) treating disorders caused by damage to brain cells, particularly by retarding or preventing brain damage in memory associated brain areas such as found in Alzheimer's and similar conditions.

These compositions may be taken as nutritional aids, for example for athletes, or for the treatment of medical conditions, particularly those associated with poor cardiac efficiency, insulin resistance and neuronal damage. The invention further provides methods of treatment and novel esters and polymers for inclusion in the compositions of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2001:44413 USPATFULL

TITLE: Therapeutic compositions

INVENTOR(S): Veech, Richard L., Rockville, MD, United States  
PATENT ASSIGNEE(S): BTG International Limited, London, United Kingdom  
(non-U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6207856	B1	20010327
APPLICATION INFO.:	US 2000-630007		20000731 (9)
RELATED APPLN. INFO.:	Division of Ser. No. US 1999-397100, filed on 16 Sep 1999 Continuation of Ser. No. WO 1997-US9805072, filed on 17 Mar 1997		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-40858P	19970317 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Reamer, James H.	
LEGAL REPRESENTATIVE:	Nixon & Vanderhye	
NUMBER OF CLAIMS:	4	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 2 Drawing Page(s)	
LINE COUNT:	1949	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 8 OF 10 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.  
on STN

TI Butyrylcholinesterase K variant on chromosome 3 q is associated with Type II diabetes in white Caucasian subjects.

AB Aims/hypothesis. To determine the association of three genes associated with Alzheimer's disease - butyrylcholinesterase (BcHE) on chromosome 3 q,  $\alpha 2$  macroglobulin ( $\alpha 2$ M) on chromosome 12p and apolipoprotein E (ApoE) on chromosome 19 q -with Type II (non-insulin-dependent) diabetes mellitus. Methods. Frequencies of BcHE K variant,  $\alpha 2$ M insertion and/or deletion polymorphism, the ApoE common polymorphisms and promoter variants at ApoE-491 and -291, were examined by fluorescent RFLP in DNA from 276 United Kingdom Prospective Diabetes Study Type II diabetic subjects and 351 non-diabetic subjects from the Diabetes In Families study. Genetic data in diabetic subjects was analysed in relation to clinical characteristics and islet function as assessed by the requirement for **insulin therapy** 6 years after randomisation.

Results. The BcHE K variant allele was more common among Type II diabetic subjects (D) than non-diabetic subjects (ND) (22.8% D vs 15.8% ND;  $p = 0.00017$ ). Subjects homozygous for the variant were more frequent in the diabetic group (5.8% D vs 2.6% ND;  $p = 0.00056$ ). The K variant allele frequency was not associated with a requirement for **insulin therapy** (29.0% insulin-requiring vs 21.8% non-insulin-requiring;  $p = 0.121$ ). There were no associations of  $\alpha 2$ M and ApoE polymorphisms or ApoE promoter variants with clinical characteristics or insulin requirement in diabetic subjects. There were differences in total cholesterol ( $p = 0.0005$ ) and LDL-cholesterol ( $p = 0.0009$ ) among non-diabetic subjects in relation to ApoE-491 genotypes.

Conclusion/interpretation. The association of the BcHE gene (3q26) with Type II diabetes could be related to an identified susceptibility locus on chromosome 3q27 but appears to be independent of islet function. The absence of diabetes-specific associations with  $\alpha 2$ M, ApoE or ApoE promoter variants suggest that these are not important in the onset of hyperglycaemia.

ACCESSION NUMBER: 2002005139 EMBASE

TITLE: Butyrylcholinesterase K variant on chromosome 3 q is associated with Type II diabetes in white Caucasian subjects.

AUTHOR: Hashim Y.; Shepherd D.; Wiltshire S.; Holman R.R.; Levy J.C.; Clark A.; Cull C.A.

CORPORATE SOURCE: C.A. Cull, Diabetes Trials Unit, Oxford Centre for Diabetes, Endocrinology and Metabolism, Woodstock Road, Oxford OX2 6HE, United Kingdom. carole.cull@dtu.ox.ac.uk

SOURCE: Diabetologia, (2001) 44/12 (2227-2230).  
 Refs: 11  
 ISSN: 0012-186X CODEN: DBTGJ

COUNTRY: Germany

DOCUMENT TYPE: Journal; Article

FILE SEGMENT: 003 Endocrinology  
 006 Internal Medicine  
 017 Public Health, Social Medicine and Epidemiology  
 022 Human Genetics  
 037 Drug Literature Index

LANGUAGE: English

SUMMARY LANGUAGE: English

L9 ANSWER 9 OF 10 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN  
 TI Apolipoprotein E genotype, islet amyloid deposition and severity of Type 2 diabetes.

AB Islet amyloid is found in 90% of patients with Type 2 (non-insulin-dependent) diabetes at post-mortem. More extensive amyloidosis is associated with decreased islet function and requirement for **insulin therapy**. Severity of cerebral amyloidosis in Alzheimer's disease (AD) is increased in subjects with the apolipoprotein E (ApoE) epsilon4 allele. To determine if ApoE genotype was associated with severity of islet amyloidosis and diabetes, samples were genotyped from 32 specimens of post-mortem pancreas and from patients classified by disease progression. DNA was extracted from blood samples from Caucasian patients diagnosed with Type 2 diabetes, at age >40 years, classified according to disease progression: group 1 on oral therapy for at least 10 years from diagnosis, (n=147) and group 2, requiring insulin within 6 years from diagnosis, (n=187). ApoE genotype was determined by restriction-fragment length polymorphism analysis. DNA in pancreatic extracts (23 diabetic; 9 non-diabetic subjects) showed no association of ApoE polymorphisms with either degree of islet amyloidosis or disease severity. The distributions of ApoE epsilon2, epsilon3 and epsilon4 were similar in both clinical patient groups and in the non-diabetic group and unrelated to progression of disease. It is unlikely that the common polymorphisms for the ApoE gene are linked to amyloid formation or progression of islet dysfunction in Type 2 diabetes.

ACCESSION NUMBER: 2003:280234 BIOSIS

DOCUMENT NUMBER: PREV200300280234

TITLE: Apolipoprotein E genotype, islet amyloid deposition and severity of Type 2 diabetes.

AUTHOR(S): Powell, D. S.; Maksoud, H.; Charge, S. B. P.; Moffitt, J. H.; Desai, M.; Da Silva Fihlo, R. L.; Hattersley, A. T.; Stratton, I. M.; Matthews, D. R.; Levy, J. C.; Clark, A. [Reprint Author]

CORPORATE SOURCE: Diabetes Research Laboratories, Oxford Centre for Diabetes Endocrinology and Metabolism, Radcliffe Infirmary, Oxford, OX2 6HE, UK  
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SOURCE: Diabetes Research and Clinical Practice, (May 2003) Vol. 60, No. 2, pp. 105-110. print.  
 ISSN: 0168-8227 (ISSN print).

DOCUMENT TYPE: Article

LANGUAGE: English

ENTRY DATE: Entered STN: 11 Jun 2003

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L9 ANSWER 10 OF 10 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

TI A case of diabetic hyperosmolar nonketotic coma with rhabdomyolysis accompanied by Alzheimer's disease.

AB A-69-year-old demented woman who had appetite loss was referred to this

hospital in June 17, 1997. She was diagnosed as having diabetes mellitus (DM) and Alzheimer's disease. Conventional **insulin therapy** was started. Her basal cortisol level and cortisol circadian rhythm were elevated (29.8 mug/dl at 07 : 00) and the postdexamethasone (1 mg) cortisol level was not completely suppressed (4.7 mug/dl). She was discharged on August 30, 1997. However, she was readmitted because of loss of consciousness on September 6, 1997. Her blood glucose level was 969 mg/dl, serum CK level was 1,137 IU/l, and serum myoglobin level was 1,784 mg/dl. Her CRP was within normal limits and urinary ketone body was negative. She was diagnosed as having hyperosmolar nonketotic coma (HNKC) with rhabdomyolysis. Transfusion of half saline and continuous administration of insulin were started, and on the fifth hospital day, her consciousness and electrolytes substantially returned to normal. CK and myoglobin levels gradually decreased. In this case, the development of HNKC was assumed to be precipitated by an acute viral infection because there were no findings of bacterial infection by diagnostic imaging techniques. The above described clinical data of this patient is suggestive of the close association between the onset of HNKC and the prolonged glucocorticoid hypersecretion under stress conditions that exists in DM when accompanied by Alzheimer's disease.

ACCESSION NUMBER: 2000:5485 BIOSIS  
DOCUMENT NUMBER: PREV200000005485  
TITLE: A case of diabetic hyperosmolar nonketotic coma with rhabdomyolysis accompanied by Alzheimer's disease.  
AUTHOR(S): Azami, Yasushi [Reprint author]  
CORPORATE SOURCE: Department of Internal Medicine, Jouhoku Hospital, Kanazawa, Japan  
SOURCE: Journal of the Japan Diabetes Society, (1999) Vol. 42, No. 6, pp. 461-465. print.  
CODEN: TONYA4. ISSN: 0021-437X.  
DOCUMENT TYPE: Article  
LANGUAGE: Japanese  
ENTRY DATE: Entered STN: 23 Dec 1999  
Last Updated on STN: 31 Dec 2001

=> d his

(FILE 'HOME' ENTERED AT 13:35:55 ON 13 FEB 2004)

FILE 'MEDLINE, USPATFULL, DGENE, EMBASE, WPIDS, BIOSIS' ENTERED AT 13:36:29 ON 13 FEB 2004

L1 92358 S ALZHEIMER DISEASE  
L2 1284 S L1 AND INSULIN  
L3 0 S INSULIN AILMENTS  
L4 18 S INSULIN ( ) DISEASE  
L5 1 S L4 AND L1  
L6 0 S GLUCOSE THERAPY AND L1  
L7 159 S GLUCOSE THERAPY  
L8 11261 S INSULIN THERAPY\  
L9 10 S L8 AND L1  
L10 0 S L7 AND L1

=> s ionic and hydrophobic compounds

L11 659 IONIC AND HYDROPHOBIC COMPOUNDS

=> s l11 and insulin

L12 191 L11 AND INSULIN

=> s insulin agonist

L13 3106 INSULIN AGONIST

=> s l13 and l12

L14 1 L13 AND L12

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L14 ANSWER 1 OF 1 USPATFULL on STN

TI 207 human secreted proteins

AB The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:258639 USPATFULL

TITLE: 207 human secreted proteins

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PRIORITY INFORMATION:	US 2000-184836P	20000224 (60)
	US 2000-193170P	20000329 (60)
	US 1997-48885P	19970606 (60)
	US 1997-49375P	19970606 (60)
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US 1997-70923P	19971218 (60)
US 1998-92921P	19980715 (60)
US 1998-94657P	19980730 (60)
US 1997-70923P	19971218 (60)
US 1998-92921P	19980715 (60)
US 1998-94657P	19980730 (60)

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EXEMPLARY CLAIM: 1  
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